

### **In the Claims**

1. (Previously presented) A method of preferentially delivering an active agent to a reservoir cell of a mammalian subject comprising:

administering to the mammalian subject a lipid-active agent complex comprising the active agent and further comprising at least one targeting ligand on the outer surface of the lipid-active agent complex that binds a group/family of markers on the surface of the reservoir cell, the reservoir cell being infected with, or susceptible to infection with an infectious agent, wherein the reservoir cell is selected from the group consisting myeloid dendritic cells, macrophages and monocytes and the targeting ligand is selected from the group consisting of fucose, polyfucose and polyfucose derivative.

2. (Previously presented) The method of Claim 1, wherein the infectious agent is a virus, bacterium, fungus or protozan.

3. -5 (Cancelled)

6. (Original) The method of Claim 2, wherein the virus is selected from the group consisting of HIV-1, HIV-2, HCV, CMV, HSV, EBV, HPV, influenza virus, and Ebola virus.

7. (Withdrawn) The method of Claim 2, wherein the bacterium is selected from the group consisting of Mycobacterium tuberculosis and Mycobacterium spec.

8. (Withdrawn) The method of Claim 2, wherein the protozoan is selected from the group consisting of Leishmania amastigotes and the discrete maturation stages of the Plasmodium life cycle.

9. (Original) The method of Claim 1, wherein the lipid-active agent complex is a liposome- active agent complex.

10. (Previously presented) The method of Claim 1, wherein the active agent is a plant lectin, an anti-viral drug, an anti-HIV drug, an anticancer drug, a cytotoxic agent, an apoptosis inhibitor, an antifungal drug, an antibacterial drug, or an immunomodulatory agent.

11. -12 (Cancelled)

13. (Withdrawn) The method of Claim 12, wherein the active agent is indinavir, saquinavir, nelfinavir, or tenofovir disoproxil fumarate.
14. (Cancelled)
15. (Original) The method of Claim 1, wherein the lipid-active agent complex further comprises one or more secondary active agents.
16. (Previously presented) The method of Claim 1, wherein the lipid-active agent complex further comprises one or more accessory factors, wherein the accessory factors is bivalent cations, co-enzymes, enzyme activators, or pH-modifying agents.
- 17.-19. (Cancelled)
20. (Withdrawn) The method of Claim 1, wherein the active agent is a small interfering RNA (siRNA).
21. (Withdrawn) The method of Claim 1, wherein the active agent is a sense or an anti-sense RNA.
22. (Withdrawn) The method of Claim 1, wherein the active agent is an expression vector suitable for dendritic cell-mediated vaccination, such as tumor vaccination.
23. (Withdrawn) The method of Claim 1, wherein the active agent is a preprocessed protein or peptide suitable for dendritic cell-mediated vaccination, such as tumor vaccination.
24. (Withdrawn) The method of Claim 10, wherein the immunomodulatory agent is an immunosuppressant or immunoactivating agent.
25. (Cancelled)
26. (Original) The method of Claim 9, wherein the active agent is encapsulated in the liposome of the liposome-active agent complex.
27. (Original) The method of Claim 1, wherein the infectious agent is susceptible to the active agent.

28. (Previously presented) The method of Claim 1, wherein the administering is by a transvascular route, a subcutaneous route, an intradermal route, a bone-marrow- directed route, an intraplacental route, an intrauterine route, intrahepatic route, an intraperitoneal route or a parenteral route.

29.-36. (Cancelled).

37. (Previously presented) The method of claim 28, wherein the administering by the intrahepatic route by infusion into the hepatic artery.

38. -49. (Cancelled)

50. (Previously presented) The method of Claim 10, wherein the plant lectin is Concanavalin A (Con-A) or myrianthus holstii lectin (MHL).

51. (Cancelled)

52. (Original) A method of preferentially delivering a plant lectin to a reservoir cell of a mammalian subject comprising:

administering to the mammalian subject a lipid-active agent complex comprising a plant lectin and further comprising at least one fucose, polyfucose, or polyfucose derivative that binds a CTL/CTLD receptor on the surface of the reservoir cell, the reservoir cell being infected with, or susceptible to infection with, an infectious agent.

53. (Previously presented) The method of claim 52, wherein the plant lectin is Concanavalin A (Con-A) or myrianthus holstii lectin (MHL).

54. (Cancelled)

55. (Withdrawn) The method of claim 52, wherein the polyfucose derivative is a fucosyl-cholesterol derivative.

56. (Original) The method of claim 53, wherein the lipid-plant lectin complex further comprises  $\text{Ca}^{2+}$  and transition-metal ions.

57. (Previously presented) The method of claim 53, wherein the MHL is a dimeric or multimeric variant of MHL.

58. (Previously presented) The method of claim 52, wherein the lipid-plant lectin complex comprises a lipid to plant lectin molar ratio between 5:1 to 7: 1.

59. (Original) The method of claim 52, wherein the lipid-plant lectin complex is between 30- 250 nm in diameter.

60. -74 (Cancelled)

75. (Withdrawn) A method for preferentially delivering an active agent to a cell with a chronic non-infectious disease comprising,

administering a lipid-active agent complex comprising the active agent and further comprising at least one targeting ligand on the outer surface of the lipid-active agent complex, wherein the targeting ligand binds a marker on the cell.

76. (Withdrawn) A method for treating HIV infected cells comprising:

administering a liposome-plant lectin complex to the HIV infected cells, wherein the outer surface of the liposome comprises a fucose derivative.

77. (Withdrawn) The method of claim 76, wherein the fucose derivative is Fuc-4C-Chol.

78. (Withdrawn) The method of claim 76, wherein the plant lectin is Con-A.

79. (Withdrawn) The method of claim 76, wherein the administering is by a subcutaneous route.

80. (Cancelled)

81. (Previously presented) A method for the intracellular delivery of an active agent to a reservoir cell comprising, administering a lipid-active agent complex to the reservoir cell, wherein the lipid-active active agent complex comprises an active agent that is encapsulated in the complex and further comprises a carbohydrate recognition domain (CRD) receptor-specific targeting ligand on the outer surface of the lipid-active agent complex, wherein the CRD receptor-specific targeting ligand is selected from fucose, polyfucose and polyfucose derivative.